'Safety'? How about we start referring to these as 'Hazards'.

A little birdie sent me an email... keep 'em coming. And way to go team.

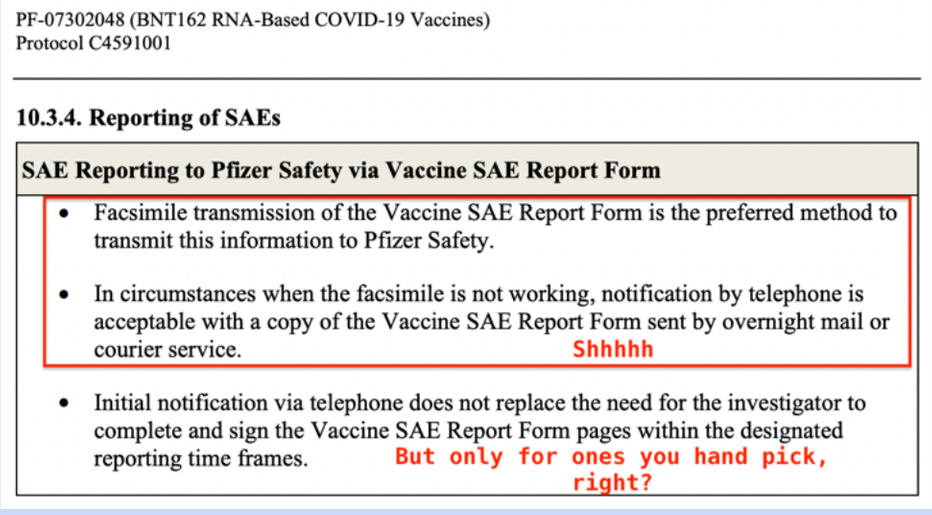
Jessica Rose
Jul 19

451 35

Today I received an email from a citizen warrior with two documents attached. By the way, I will post everything you guys send, no matter how 'sensitive'. We paid for this program and these products, and we not only deserve to know what's being injected into our species, but are **entitled** to know. I can't believe that what I just wrote is deemed 'controversial'.

The two documents are Pfizer 'safety' documents readily available online entitled Pfizer Safety Data Sheet and A phase 1/2/3, placebo-controlled, randomized, observer-blind, dose-finding study to evaluate the safety, tolerability, immunogenicity, and efficacy of SARS-nCoV-2 RNA vaccine candidates against COVID-19 in healthy individuals.

The emailer referred me to page 131 in the latter document where something peculiar is documented.



PF-07302048 (BNT162 RNA-Based COVID-19 Vaccines)
Protocol C4591001

10.3.4. Reporting of SAEs

SAE Reporting to Pfizer Safety via Vaccine SAE Report Form

- Facsimile transmission of the Vaccine SAE Report Form is the preferred method to transmit this information to Pfizer Safety.
- In circumstances when the facsimile is not working, notification by telephone is acceptable with a copy of the Vaccine SAE Report Form sent by overnight mail or courier service. Shhhhh
- Initial notification via telephone does not replace the need for the investigator to complete and sign the Vaccine SAE Report Form pages within the designated reporting time frames. But only for ones you hand pick, right?

Figure 1: file:///Users/jessicarose/Downloads/14c84539-487f-4552-bac7-f412bfaad5c2.pdf. Page 131.

All SAEs (severe adverse events) were to be reported by fax or phone. There is only one reason to exclude initial notification by formal written documentation for an SAE: to ensure that if a report needs to be 'erased', it can be. My emailer put it this way:

> Why do they request all SAEs to be transmitted via ONLY facsimile, telephone, and hard-copy mail? Because, in my humble opinion, there then is no electronic record/trail.

I think my emailer is 100% correct. If I was in charge of SAE reports for a questionable 'clinical trial' and I wanted to vet the reports such that if, say, there were a lot of deaths being reported in temporal proximity to injecting my tremendously safe products into people, then I would make damned sure that those reports came privately and securely to me (without a paper trail) so that I would never have to fess up to their existence if I didn't want to, and thus I could always say that my creepy injectable products were safe and effective.

On page 128, they also state the following:

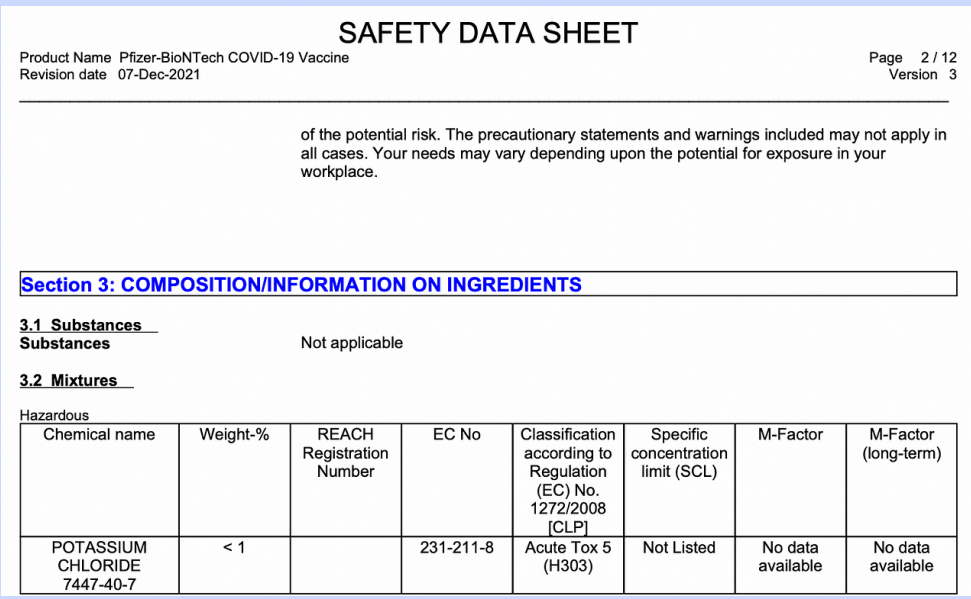
> It should be noted that the Vaccine SAE Report Form for reporting of SAE information is not the same as the AE page of the CRF.

They do write that in addition to that phone call, you have to do your paperwork, but, to be litigious, their bestie CDC also claim that VAERS reports are compulsory to file is an AE is even suspected, while simultaneously creating threatening environments in medical establishments for doctors who do want to file to encourage them not to.

And by the way, they are meant to be doing causality assessments as well.

> For each AE/SAE, the investigator must document in the medical notes that he/she has reviewed the AE/SAE and has provided an assessment of causality. There may be situations in which an SAE has occurred and the investigator has minimal information to include in the initial report to the sponsor. However, it is very important that the investigator always make an assessment of causality for every event before the initial transmission of the SAE data to the sponsor.

Here's the second piece of information from the former document. If you refer to the Safety Data Sheet on page 2, you'll find something called Potassium chloride in the Hazardous ingredients table. In column 5, you'll note a classification of Acute Tox 5 (H303).



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of the potential risk. The precautionary statements and warnings included may not apply in all cases. Your needs may vary depending upon the potential for exposure in your workplace.

Section 3: COMPOSITION/INFORMATION ON INGREDIENTS

3.1 Substances
Substances Not applicable

3.2 Mixtures

Hazardous

Chemical name	Weight-%	REACH Registration Number	EC No	Classification according to Regulation (EC) No. 1272/2008 [CLP]	Specific concentration limit (SCL)	M-Factor	M-Factor (long-term)
POTASSIUM CHLORIDE 7447-40-7	< 1		231-211-8	Acute Tox 5 (H303)	Not Listed	No data available	No data available

Figure 2: https://safetydatasheets.pfizer.com/DirectDocumentDownloader/Document?prd=PF00092~~PDF~~MTR~~PFEM~~EN. Page 2.

Potassium chloride is a mineral found in many foods and is essential for proper functioning of the beating of the heart. It can be taken as a therapeutic to prevent, or aid in, elevation of low serum levels of potassium in the context of a condition called hypokalemia. Side effects of excessive potassium include uneven heartbeat, muscle weakness or limp feeling, severe stomach pain, and numbness or tingling in your hands, feet, or mouth[1] and thus ingestion of this medication should be monitored carefully. In concentrated injected form[2], its intended use is for maintenance of serum potassium.

Notice the list of adverse reactions associated with administration of highly concentrated IV potassium chloride in the following Table found on page 6 of the FDA document found here.

ADVERSE REACTIONS

The following adverse reactions associated with the use of Potassium Chloride Injection were identified in postmarketing reports. Because these reactions were reported voluntarily from a population of uncertain size, it is not always possible to estimate their frequency, reliably, or to establish a causal relationship to drug exposure

Immune system disorders: Hypersensitivity, as manifested by rash and angioedema

Metabolism and nutrition disorders: Hyperkalemia

Cardiac disorders: Cardiac arrest*, asystole*, ventricular fibrillation*, bradycardia

*as a manifestation of rapid intravenous administration and/or of hyperkalemia

Respiratory, Thoracic, and Mediastinal Disorders: Dyspnea

General disorders and administration site conditions: Chest pain, infusion site thrombosis, infusion site phlebitis, infusion site erythema,- infusion site swelling, infusion site pain, infusion site irritation, and/or a burning sensation.

The following adverse reactions were reported in association with extravasation: Skin necrosis, skin ulcer, soft tissue necrosis, muscle necrosis, nerve injury, tendon injury, and vascular injury.

Figure 3:
https://www.accessdata.fda.gov/drugsatfda_docs/label/2017/019904s014lbl.pdf.
Page 6.

You're not meant to administer the IV concentrated form of potassium chloride if one has a cardiac disorder.

Question: What does this mean for the children who are diagnosed with myocarditis in temporal association with these shots?

They have a diagnosed cardiac disorder. Injection of potassium chloride is contraindicated with diagnosed cardiac disorders. Therefore, knowingly injecting someone with a contraindicated substance is medical malfeasance and that administrator is personally liable.

The dose 2 effect observed in myocarditis in children may be precisely this: the primary cardiac damage exacerbated by the secondary shot due to the presence of potassium chloride. To me, it would be essential that the concentration of potassium chloride was known. As it stands, according to the Pfizer Safety Data Sheet, the 'Specific concentration limit (SCL)' is 'Not Listed'.

They also have Potassium phosphate listed in the NonHazardous ingredients and injection of potassium phosphate is contraindicated with hyperkalemia.

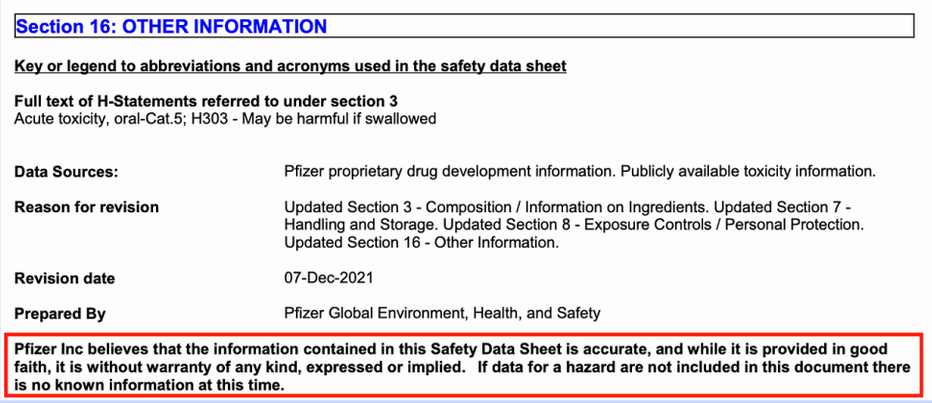
I looked H303 up in the GHS Classification on Pubchem.

H300	Fatal if swallowed	Acute toxicity, oral	Category 1, 2	Div 2.3, 6.1		Danger	P264, P270	P301+P316, P321, P330	P405	P501
H301	Toxic if swallowed	Acute toxicity, oral	Category 3	Div 2.3, 6.1		Danger	P264, P270	P301+P316, P321, P330	P405	P501
H302	Harmful if swallowed	Acute toxicity, oral	Category 4			Warning	P264, P270	P301+P317, P330		P501
H303	May be harmful if swallowed	Acute toxicity, oral	Category 5		None	Warning		P301+P317		
H304	May be fatal if swallowed and enters airways	Aspiration hazard	Category 1			Danger		P301+P316, P331	P405	P501
H305	May be fatal if swallowed and enters airways	Aspiration hazard	Category 2			Warning		P301+P316, P331	P405	P501
H310	Fatal in contact with skin	Acute toxicity, dermal	Category 1, 2	Div 2.3, 6.1		Danger	P262, P264, P270, P280	P302+P352, P316, P321, P361+P364	P405	P501
H311	Toxic in contact with skin	Acute toxicity, dermal	Category 3	Div 2.3, 6.1		Danger	P280	P302+P352, P316, P321, P361+P364	P405	P501

Figure 4: https://pubchem.ncbi.nlm.nih.gov/ghs/.

The reason why data is missing in a classification of toxicity is usually because the data has not been collected to date and therefore nothing is actually known about the toxicity at the time of publication of the document. “No data available.” Funny that. They put this stuff into 5 billion people before the toxicity data was available and said screw the environment. But, climate change.

The classification of this chemical starts with an ‘H’ and all of the Hs look pretty bad according to the Pictograms. It says that it’s bad if you eat it but, there’s no Pictogram associated with it yet as shown in Figure 4. This to me means, they don’t know how to classify it yet. There is also a warning attached to it.



Section 16: OTHER INFORMATION

Key or legend to abbreviations and acronyms used in the safety data sheet

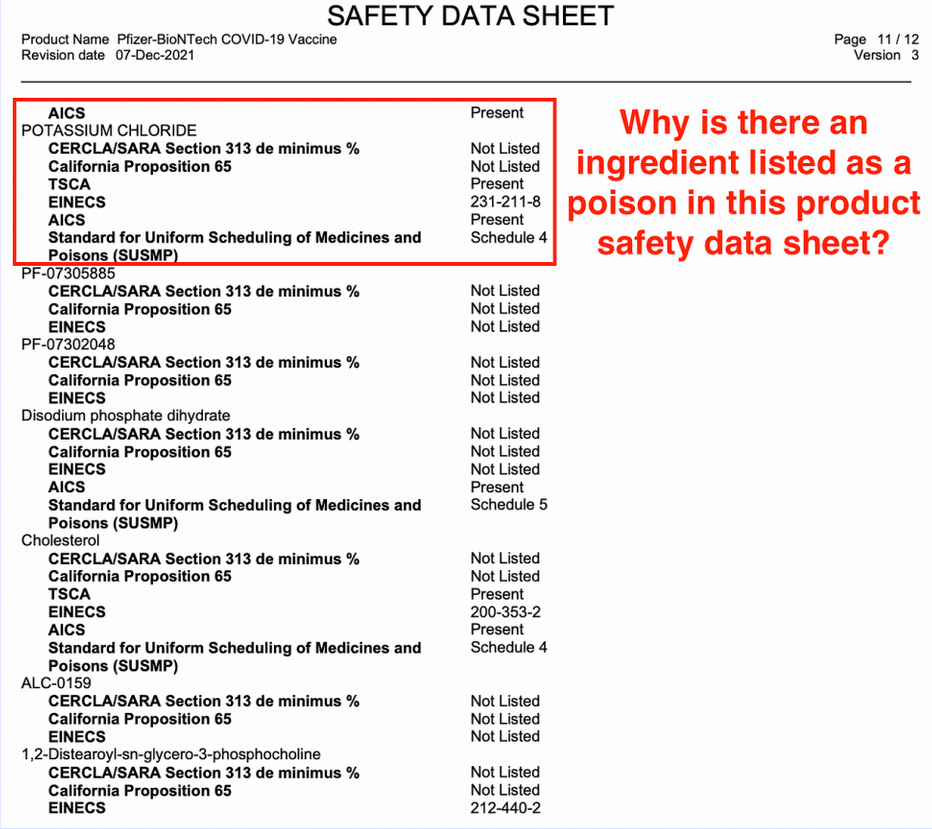
Full text of H-Statements referred to under section 3
Acute toxicity, oral-Cat.5; H303 - May be harmful if swallowed

Data Sources:	Pfizer proprietary drug development information. Publicly available toxicity information.
Reason for revision	Updated Section 3 - Composition / Information on Ingredients. Updated Section 7 - Handling and Storage. Updated Section 8 - Exposure Controls / Personal Protection. Updated Section 16 - Other Information.
Revision date	07-Dec-2021
Prepared By	Pfizer Global Environment, Health, and Safety

Pfizer Inc believes that the information contained in this Safety Data Sheet is accurate, and while it is provided in good faith, it is without warranty of any kind, expressed or implied. If data for a hazard are not included in this document there is no known information at this time.

Figure 5:
https://safetydatasheets.pfizer.com/DirectDocumentDownloader/Document?prd=PF00092~~PDF~~MTR~~PFEM~~EN. Page 12.

Pfizer ‘in good faith’? Are you kidding me? Pfizer has paid out the most in health care fraud settlements in ***history***[3] “to resolve criminal and civil liability arising from the illegal promotion of certain pharmaceutical products” according to the United States Justice Department. THEY’VE ALREADY BEEN CAUGHT RED-HANDED HARMING THE AMERICAN PUBLIC! What more do people want? When they claim that the information they provide to the public is ‘in good faith’, I say, alrighttyyyyyy thennnnnnnnn. And when they subsequently claim that un-included hazard data is ‘not known at this time’. I say, alroooioightyyyyyyy thennnnnnnn. Not known to us suckers anyway.



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AICS	Present
POTASSIUM CHLORIDE	
CERCLA/SARA Section 313 de minimus %	Not Listed
California Proposition 65	Not Listed
TSCA	Present
EINECS	231-211-8
AICS	Present
Standard for Uniform Scheduling of Medicines and Poisons (SUSMP)	Schedule 4
PF-07305885	
CERCLA/SARA Section 313 de minimus %	Not Listed
California Proposition 65	Not Listed
EINECS	Not Listed
PF-07302048	
CERCLA/SARA Section 313 de minimus %	Not Listed
California Proposition 65	Not Listed
EINECS	Not Listed
Disodium phosphate dihydrate	
CERCLA/SARA Section 313 de minimus %	Not Listed
California Proposition 65	Not Listed
EINECS	Not Listed
AICS	Present
Standard for Uniform Scheduling of Medicines and Poisons (SUSMP)	Schedule 5
Cholesterol	
CERCLA/SARA Section 313 de minimus %	Not Listed
California Proposition 65	Not Listed
TSCA	Present
EINECS	200-353-2
AICS	Present
Standard for Uniform Scheduling of Medicines and Poisons (SUSMP)	Schedule 4
ALC-0159	
CERCLA/SARA Section 313 de minimus %	Not Listed
California Proposition 65	Not Listed
EINECS	Not Listed
1,2-Distearoyl-sn-glycero-3-phosphocholine	
CERCLA/SARA Section 313 de minimus %	Not Listed
California Proposition 65	Not Listed
EINECS	212-440-2

Why is there an ingredient listed as a poison in this product safety data sheet?

Figure 6:
https://safetydatasheets.pfizer.com/DirectDocumentDownloader/Document?prd=PF00092~~PDF~~MTR~~PFEM~~EN. Page 11.

I may not be up-to-date on what a poison is, but I would rather not get injected with something with the word poison associated with it.

How about you?

Thank you to my emailer.

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1 https://www.drugs.com/potassium_chloride.html

2 https://www.accessdata.fda.gov/drugsatfda_docs/label/2017/019904s014lbl.pdf

3 https://www.justice.gov/opa/pr/justice-department-announces-largest-health-care-fraud-settlement-its-history

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451 35

35 Comments

Write a comment...

Hubris Jul 20

"Pfizer has paid out the most in health care fraud settlements in history3 “to resolve criminal and civil liability arising from the illegal promotion of certain pharmaceutical products” according to the United States Justice Department. "

Honestly, NO fine can do justice; they need to be disbanded. They were awarded the same status as the most crooked banks: "too big to fail"... Says who?

63ReplyCollapse

5 replies by Jessica Rose and others

Pete Jul 20 Liked by Jessica Rose

The last time I checked, safety had a literal meaning consistent with the substance in some road signs.

Stop means stop, that’s the same sign that is printed on all official documents requesting immediate discontinuation of manufacture of toxic fertilisers, medical substances with unknown future effects, vehicle tyres made with addition of recycled PET bottles, construction materials with asbestos, brake pads that beautifully spread completely untested powdered residues all over the world, fluorine-enriched industrial waste converted into dental care products, and so on. Dream on.

Speed limit applies to the performance of the vehicle (measured with the status of the driver), and not to road conditions. That one is obvious. Limit means “test it now”. “Straight ahead only” is the campaign logo for the new generation of injectable warpspeeders. Roadway condition signs may be a little confusing. Bump - whom? Soft shoulder - a massage service? Rough road, oh, yes, with hangover, always.

Products of the chemical industry (occasionally called pharmaceuticals) is another world. For the starters, you may check regular liquid soap. It has from a few to about 20 ingredients. They are miraculously kept “isolated” - no chemical reactions between and among them (really?). If this is true, it means that liquid soap retains the chemical and biological properties of individual components. Like caustic. Or irritating to skin. Or not tested for carcinogenic impact. Check in the databases, yes, we are using this stuff to “clean” our "dirty" hands. And more. But... what if they are chemically reacting 24/7? Locked in a no-escape room (escape room?), what else would they do? Did anyone test it? For how long? In sunshine?

Safety signals - that one is great. They should be more accurately called “reports of drug-related conditions possibly leading to aggravation of health”. With emphasis on “grave” like in the “dead end” road sign.

These cute interludes in the responsible work of the author-ities author-ising products that they have never seen or used are nothing like human safety or like warning signals. They are not about your safety. It's all about the safety of the manufacturer and the enforcement chain downline. And, yes, they are signalling potential problem areas for the stakeholders.

Who told you to interpret this as "risk to human health"?

The truth is out there, we only need to change the way we are looking at it and the way we are interpreting it.

♡ 30ReplyCollapse

33 more comments...

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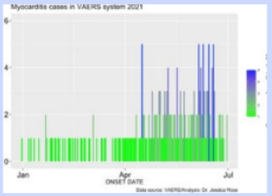
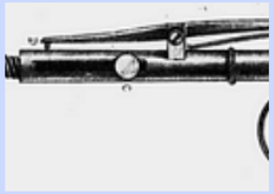


A Report on Myocarditis Adverse Events in the U.S. Vaccine Adverse Events Reporting System (VAERS) in Association with COVID-19 Injectable...

Jessica Rose PhD, MSc, BSc and Peter A. McCullough MD, MPH

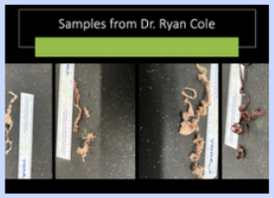
JESSICA ROSENOV 2, 2021 ♡1,206 💬143



This is one of the emails I received the other day. I get hundreds daily, and I am hearing you all.

This particular note spoke loudly to me and this lovely person gave me permission to share her words.

JESSICA ROSEJUL 17 ♡1,403 💬125



Rewrite: Let's tag team this until everybody understands

The modified spike protein is dangerous and for very specific reasons.

JESSICA ROSEJUN 13 ♡606 💬147

See all ›

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